U.S. Patent Application No. 09/888,008 Amendment C January 24, 2005

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Amended Claims

1. (currently amended) A method of determining enzyme activity, the method comprising:

contacting a compound selected from the group consisting of enzymes, enzyme fragments and abzymes with a labeled substrate having a label to form a differentially-charged product;

selectively coupling either the substrate or the differentially-charged product to an ion-exchange resin thereby substantially separating the amount of substrate from the differentially-charged product; and

detecting and quantitating the label to determine the amount of how much substrate is remaining or how much differentially-charged product is formed, wherein the steps of coupling and detecting are performed sequentially without removing the substrate or product that is not coupled to the resin.

2. (currently amended) A method of determining enzyme activity, the method comprising:

contacting a compound selected from the group consisting of enzymes, enzyme fragments and abzymes with a labeled substrate having a label thereby effecting the conversion converting of the substrate to a differentially-charged product;

stopping the conversion before all of the substrate present has been converted to the differentially-charged product;

selectively coupling either the substrate or the differentially-charged product to an ion-exchange resin thereby substantially separating the substrate from the differentially-charged product in a single step; and

detecting and quantitating the label to determine the amount of how much substrate is remaining or how much differentially-charged product is formed;

wherein the steps of coupling and detecting are performed sequentially without removing the substrate or product that is not coupled to the resin.

3. (original) The method of claim 1 or 2 wherein the product is bound to the resin.

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- 4. (original) The method of claim 1 or 2 wherein the substrate is bound to the resin.
- 5. (original) The method of claim 1 or 2 wherein the product or substrate measured is coupled to the resin.
- 6. (original) The method of claim 1 or 2 wherein the product or substrate measured is in solution.
- 7. (previously presented) The method of claim 1 or 2 wherein the enzyme is selected from the group consisting of a kinase, a transferase and a synthase.
- 8. (original) The method of claim 1 or 2 wherein said method is conducted in a multiple-well format.
- 9. (original) The method of claim 8 wherein the format comprises at least about 96 wells.

Claim 10. (canceled)

- 11. (previously presented) The method of claim 1 or 2 wherein said method is conducted in a microchip.
- 12. (previously presented) The method of claim 1 or 2 wherein said enzyme is selected from the group consisting of glutamine fructose-6-phosphate amidotransferase (GFAT), Nitric Oxide Synthase, Methionine Aminopeptidase, Asparagine Synthetase (Asn Syn), PFK, p38 kinase, I-kappa kinase 1, I-kappa kinase 2, TBK 1, MAPKAP 2, galactosyl transferase (GTase), O-n-acetylglucosamine transferase (OGTase), and Cyclooxygenase.

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13. (currently amended) A method for identifying a molecule, compound, or composition that affects the activity of an enzyme, the method comprising:

contacting the enzyme with a test sample comprising a molecule, compound, or composition;

contacting the enzyme with a labeled substrate having a label to form a differentially-charged product;

selectively coupling either the substrate or the differentially-charged product to an ion-exchange resin thereby substantially separating the substrate from the differentially-charged product;

detecting and quantitating the label to determine the amount of how much substrate is remaining or how much differentially-charged product is formed; and

comparing the amount of substrate remaining or differentially-charged product formed with a control, wherein the steps of coupling and detecting are performed sequentially without removing the substrate or product that is not coupled to the resin.

- 14. (previously presented) The method of claim 13 wherein said enzyme is selected from the group consisting of glutamine fructose-6-phosphate amidotransferase (GFAT), Nitric Oxide Synthase, Methionine Aminopeptidase, Asparagine Synthetase (Asn Syn), PFK, p38 kinase, I-kappa kinase 1, I-kappa kinase 2, TBK1, MAPKAP 2, galactosyl transferase (GTase), O-n-acetylglucosamine transferase (OGTase), and Cyclooxygenase.
- 15. (previously presented) The method of claim 13 wherein the control is an isozyme and the method is used to identify a compound or composition that preferentially or specifically affects an enzyme over its isozyme.

Claims 16-26. (canceled)